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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/035,637	11/07/2001	Yashwant M. Deo	CDJ-166CP	4452
959	7590	05/23/2006	EXAMINER	
LAHIVE & COCKFIELD 28 STATE STREET BOSTON, MA 02109			EWOLDT, GERALD R	
			ART UNIT	PAPER NUMBER
			1644	
DATE MAILED: 05/23/2006				

Please find below and/or attached an Office communication concerning this application or proceeding.

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DETAILED ACTION

1. A request for continued examination (RCE) under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed 3/30/06 in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's amendments, remarks, and IDS, filed 3/30/06, have been entered.

2. Claims 42-46 stand withdrawn from further consideration by the examiner, 37 CFR 1.142(b), as being drawn to non-elected inventions.

Claims 27-35, 38, 39, 51, and newly added Claim 52 read on the elected invention and are being acted upon.

3. In view of the instant amendment and remarks, deleting the term "conservative sequence modifications" and citing additional support for the molecular conjugate of the instant claims, the previous rejections under the first paragraph of 35 U.S.C. 112 have been withdrawn.

4. The title and abstract are objected to because they do not accurately describe the claimed invention. Correction is required.

5. This application contains sequence disclosures that are encompassed by the definitions for nucleotide and/or amino acid sequences set forth in 37 C.F.R. 1.821(a)(1) and (a)(2). However, this application fails to comply with the requirements of 37 C.F.R. 1.821-1.825 for the reason(s) set forth on the attached Notice To Comply With Requirements For Patent Applications Containing Nucleotide Sequence And/Or Amino Acid Sequence Disclosures. Applicant must comply with the requirements of the sequence rules (37 CFR 1.821 - 1.825) before the application can be examined under 35 U.S.C. 131 and 132. Specifically, the sequences in Figure 13 must be identified by SEQ ID NO:. Additionally, upon the addition of Inventors Trembl and Endres, the inventorship of the Sequence Listing and CFR is now incorrect.

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6. The instant application claims priority to Application Nos. 09/851,614, 60/203,126, and 60/230,739. A review of the '614 application shows that the molecular conjugate of said application is not the molecular conjugate of the instant claims. The conjugate of the '614 application, as recited in Claims 1 and 5, is "specific" for DCs; the conjugate of the instant claims does not include this limitation. Additionally, the conjugate of the '614 applications comprises additional limitations, also set forth in claim 1:

- a) a binding affinity constant to a dendritic cell of at least about 10^7 M^{-1} ;
- b) the ability to opsonize a dendritic cell;
- c) the ability to internalize after binding to dendritic cells;
- or
- d) the ability to activate dendritic cells.

Accordingly, priority to the '614 application is denied. The priority date of the instant application is its filing date, 11/07/01

7. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

8. Claims 27-30, 32, 33, 38, 39, 51, and newly added Claim 52 stand/are rejected under 35 U.S.C. 103(a) as being unpatentable over U.S. Patent No. 5,922,845 (IDS) in view of Tuting et al. (1998, of record) and Sallusto et al. (1995).

As set forth previously, The '845 patent teaches a molecular conjugate comprising an antibody that binds to dendritic cells (DCs) (Fc α R) and an antigen, wherein said antigen comprises a component of a pathogen or a tumor (cancer) antigen (see column 3, lines 49-59). The reference further teaches the conjugate comprising a single chain antibody (see column 3, line 63), a pharmaceutically acceptable carrier (see column 4, line 32), and an adjuvant (see column 21, line 54). The reference further teaches that the molecular conjugates of the reference can be used to "harness the capabilities of white blood cells", e.g., phagocytosis, for "enhancing the attack of these cells against cancer cells, cells of infectious microorganisms, and cells infected with pathogens".

The reference teaching differs from the claimed invention only in that it does not teach a molecular conjugate comprising an antibody that binds to a human macrophage mannose receptor and the Pmel-17 tumor antigen.

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Tuting et al. teaches that Pmel-17 is one of several well known melanoma antigen (see particularly page 1140, column 1).

Sallusto et al. teaches that the human mannose macrophage receptor (which would be encoded by SEQ ID NO:7) can be employed for the uptake of antigen by DCs for presentation of said antigen to T cells (see particularly Abstract; page 392, column 1; and Figure 4).

It would have been *prima facie* obvious to one of ordinary skill in the art at the time the invention to produce a molecular conjugate comprising an antibody that binds to white blood cells (which would include DCs) and tumor antigen, as taught by the '845 patent, employing an antibody that binds the human macrophage mannose receptor and Pmel-17 as the antigen. One of ordinary skill in the art at the time the invention was made would have been motivated to employ an antibody that binds the human macrophage mannose receptor because it is more APC-specific than is the Fc α R of the '845 patent, thus allowing for more efficient antigen uptake by APCs and more efficient antigen presentation to T cells. One of ordinary skill in the art at the time the invention was made would have been motivated to employ any of the well known tumor antigens in an anti-cancer therapeutic agent, such as Pmel-17 as taught by Tuting et al., because of their availability and previous characterization.

Applicant's arguments, filed 3/30/06, have been fully considered but they are not persuasive. Applicant argues that mAb bind targets with high affinity and such antibodies would not have been thought suitable for targeting antigens to the mannose receptor because the antigen-antibody complex would fail to dissociate upon internalization.

First note that Applicant has provided no evidence that the skilled artisan would not have expected the antigen-antibody complex to dissociate upon internalization, but regardless, the skilled artisan would have simply selected an antibody with reduced binding affinity if this were a concern.

Applicant argues, "Sallusto et al. make a clear distinction between mannose receptor mediated-endocytosis of ligands and the internalization of Fc receptors and their ligands; the latter of which results in delivery to lysosomes and the degradation of both the ligand and the receptor. Accordingly, one of ordinary skill would not have been motivated to have used antibodies to target antigens to dendritic cells since such antibodies would have been expected also to bind the Fc receptors expressed on these cells, thus, resulting in degradation of the ligand and the receptor".

Again the arguments comprises only an attorney's perception of a potential problem. Regardless, the skilled artisan could

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have simply overcome this potential problem by using more conjugate or selecting an antibody of appropriate binding affinity, i.e., an affinity slightly higher than the affinity of the FcR for the Fc region of an antibody.

Applicant discusses macropinocytosis and speculates that "such mechanisms ... would not have been thought suitable for antibody-based vaccines".

Again, Applicant provides only an attorney's speculation.

Applicant cites Ramakrishna et al. (2004) wherein the use of an antibody to target antigens to the mannose receptor, i.e., the method of the instant claims is taught. Interestingly, it is noted that the lead Inventor, Yashwant Deo, is not included as an author of the paper, but the remaining Inventors, Keler, Trembl, and Endres are. Regardless, a review of the work shows that the authors never indicate that the construction and use of a molecular conjugate for the targeting of antigens to the mannose receptor is anything other than routine. They do not indicate anything unexpected, surprising, nor particularly difficult in producing a functional conjugate. Indeed, they indicate that they are simply employing the same rationale used by Steinman et al. wherein the related DEC-205 receptor was targeted for the introduction of antigen into DCs.

9. The following is a new grounds of rejection.

10. Claims 27-30, 32, 33, 38, 39, 51, and newly added Claim 52 are rejected under 35 U.S.C. 103(a) as being unpatentable over U.S. Patent Application Publication No. 2002/0187131 (IDS) in view of U.S. Patent No. 5,922,845 (IDS), and Tuting et al. (1998), all of record.

The '131 application discloses a molecular conjugate (including recombinant and chemically conjugated), comprising an antibody (including a single chain antibody) and an antigen (including tumor or pathogen antigens), see particularly paragraphs 19, 20, 43, and 46.

The reference teaching differs from the claimed invention only in that it does not teach a molecular conjugate comprising a human monoclonal antibody and the Pmel-17 tumor antigen.

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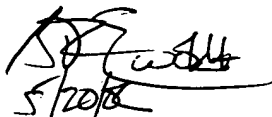
Tuting et al. and the '845 patent have been discussed above. The '845 patent further teaches the value of molecular conjugates comprising human monoclonal antibodies including the elimination of a HAMA (human anti-mouse antibody) response, see particularly column 8, line 33 - column 9, line 33).

It would have been *prima facie* obvious to one of ordinary skill in the art at the time the invention to produce the molecular conjugate of the '131 application comprising the Pmel-17 tumor antigen and a human monoclonal antibody. One of ordinary skill in the art at the time the invention was made would have been motivated to employ the Pmel-17 as taught by Tuting et al., because of its availability and previous characterization, and a human monoclonal antibody, as taught by the '845 patent for the elimination of a HAMA response.

11. Claim 31 is allowed. Claims 34 and 35 would be allowable if recited in independent form.

12. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Dr. Gerald Ewoldt whose telephone number is (571) 272-0843. The examiner can normally be reached Monday through Thursday from 7:30 am to 5:30 pm. A message may be left on the examiner's voice mail service. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Christina Chan can be reached on (571) 272-0841.

13. **Please Note:** Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).


5/20/12
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